

SENNALAX- sennosides tablet
Marc Glassman, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

371 - Senna-Lax

Active ingredient

Sennosides 8.6 mg

Purpose

Laxative

Uses

- for relief of occasional constipation (irregularity)
- this product produces a bowel movement in 6 to 12 hours

Warnings

Do not use

laxative products for more than one week unless directed by a doctor

Ask a doctor before use if you have

- stomach pain
- nausea
- vomiting
- noticed a sudden change in bowel habits that continues over a period of 2 weeks

Stop use and ask a doctor if

- rectal bleeding or failure to have a bowel movement occur after use of a laxative. These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children

In case of accidental overdose, contact a doctor or Poison Control Center (1-800-222-1222) right away.

Directions

- take preferably at bedtime or as directed by a doctor

Age	Starting dosage	Maximum dosage
Adults and children over 12 years	2 tablets once a day	4 tablets twice a day
Children 6 to under 12 years	1 tablet once a day	2 tablets twice a day
Children 2 to under 6 years	1/2 tablet once a day	1 tablet twice a day
Children under 2 years	consult a doctor	consult a doctor

Other information

■ store at 20°C-25°C (68°F-77°F)

- do not use if safety seal under cap is torn or missing
- **Calcium content:** 50 mg per tablet

Inactive ingredients

croscarmellose sodium, dibasic calcium phosphate, magnesium stearate, microcrystalline cellulose, silicon dioxide

Principal Display Panel

Senna-Lax

Senna-Lax Label
68998-373-01

†Compare to the Active Ingredient in Senokot®

Marc's (logo)

Senna-Lax

NATURAL VEGETABLE LAXATIVE

Gently Relieves Constipation

100 Tablets

SENNALAX					
sennosides tablet					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68998-373		
Route of Administration	ORAL				
Active Ingredient/Active Moiety					
Ingredient Name		Basis of Strength	Strength		
SENNOSIDES (UNII: 3FYP5M0IJX) (SENNOSIDES - UNII:3FYP5M0IJX)		SENNOSIDES	8.6 mg		
Inactive Ingredients					
Ingredient Name			Strength		
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)					
MAGNESIUM STEARATE (UNII: 70097M6I30)					
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)					
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)					
Product Characteristics					
Color	brown	Score	no score		
Shape	ROUND	Size	8mm		
Flavor		Imprint Code	AZ;217		
Contains					

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68998-373-01	1 in 1 CARTON	07/01/2014	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	07/01/2014	

Labeler - Marc Glassman, Inc. (094487477)

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Marc Glassman, Inc.